

**INFORMATION  
DISCLOSURE  
STATEMENT**

Atty. Docket No.: 249.00020101

Serial No.: 09/970,318

Applicant(s): Cosgrove

Confirmation No.: 1885

Filing Date: October 3, 2001

Group: 1641

**U.S. PATENT DOCUMENTS**

Document Number	Date	Name	Class	Subclass	Filing Date If Appropriate
none					

**FOREIGN PATENT DOCUMENTS**

Examiner Initial	Document Number	Date	Country	Class	Subclass	Translation	
						Yes	No
	none						

**OTHER DOCUMENTS (Including Authors, Title, Date, Pertinent Papers, etc.)**

Examiner Initial	Document Description
MY	Adato et al., "Three novel mutations and twelve polymorphisms identified in the USH2A gene in Israeli USH2 families," <i>Hum Mutat. (Mutation in Brief)</i> ; published online for subscribers 2000 Mar 22 (6 pgs.).
MY	Adato et al., "Three novel mutations and twelve polymorphisms identified in the USH2A gene in Israeli USH2 families," <i>Hum Mutat.</i> 2000 Apr;15(4):abst. 388.
MY	Barkalow et al., "Localization of the major heparin-binding site in fibronectin," <i>J Biol Chem.</i> 1991 Apr 25;266(12):7812-8.
MY	Beck et al., "Structure and function of laminin: anatomy of a multidomain glycoprotein," <i>FASEB J.</i> 1990 Feb 1;4(2):148-60.
MY	Bork et al., "Structure and distribution of modules in extracellular proteins," <i>Q Rev Biophys.</i> 1996 May;29(2):119-67.
MY	Boughman et al., "Usher syndrome: definition and estimate of prevalence from two high-risk populations," <i>J Chronic Dis.</i> 1983;36(8):595-603.
MY	Bowditch et al., "Identification of a novel integrin binding site in fibronectin. Differential utilization by $\beta 3$ integrins," <i>J Biol Chem.</i> 1994 Apr 8;269(14):10856-63.

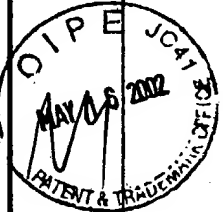
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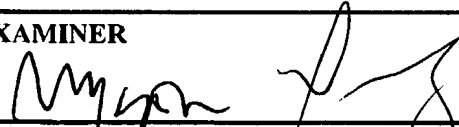
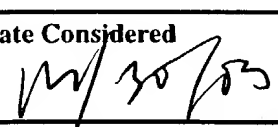
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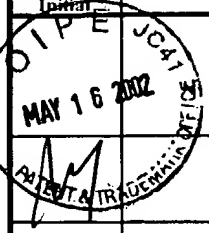
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	Brookhouser, Patrick E., "Center for Hearing Loss in Children," Grant Abstract, Grant Number 5P60DC000982-10 [online]. National Institutes of Health, National Institute on Deafness and other Disorders of Communication, project dates 1990-09-30 to 2001-08-31. Retrieved from the Internet 2002-03-21. URL: < <a href="http://commons.cit.nih.gov/crisp3/CRISP_LIB.getdoc?textkey=6055813&amp;p_grant_num=5P60DC000982-10&amp;p_query=&amp;ticket=388948&amp;p_audit_session_id=3084290&amp;p_keywords=&gt;">http://commons.cit.nih.gov/crisp3/CRISP_LIB.getdoc?textkey=6055813&amp;p_grant_num=5P60DC000982-10&amp;p_query=&amp;ticket=388948&amp;p_audit_session_id=3084290&amp;p_keywords=&gt;</a> >; 2 pages.
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	Kimberling et al., "Gene mapping of Usher syndrome type IIa: localization of the gene to a 2.1-cM segment on chromosome 1q41," <i>Am J Hum Genet.</i> 1995 Jan; 56(1):216-23.
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M	National Center for Biotechnology Information, National Library of Medicine, National Institutes of Health, GenBank Locus HSUSH2A01, Accession No. AF091873, "Homo sapiens Usher syndrome type IIa protein gene, exons 1 and 2," [online]. Bethesda, MD [retrieved on 2002-04-05]. Retrieved from the Internet: <URL:http://www.ncbi.nlm.nih.gov/entrez/viewer.fcgi?query_key=4&db=nucleotide&page=0&dispmax=20&WebEnv=Wg%3Cn_FDG%5DE%60%3E%3D%3C%5DPGDJc_gTB%5EjbFkl%3C_JEH%3Dzcc%3EF%5EFJdTI%3D%3F%3C&WebEnvRq=1>; 3 pgs.

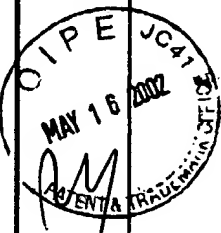
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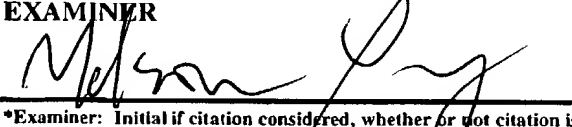
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	National Center for Biotechnology Information, National Library of Medicine, National Institutes of Health, GenBank Locus HSUSH2A17, Accession No. AF091889, "Homo sapiens Usher syndrome type IIa protein gene, exon 21 and complete cds," [online]. Bethesda, MD [retrieved on 2002-04-05]. Retrieved from the Internet: <URL: <a href="http://www.ncbi.nlm.nih.gov/entrez/viewer.fcgi?query_key=6&amp;db=nucleotide&amp;page=0&amp;dispmax=20&amp;WebEnv=C%5B%60%3CIgCAA%5C%5Ee%3D%3FHCK%3CCDB%3CdlEjhCdbaiH%3F%40AARqjH%5EGe%5E%3EkDFH&amp;WebEnvRq=1">http://www.ncbi.nlm.nih.gov/entrez/viewer.fcgi?query_key=6&amp;db=nucleotide&amp;page=0&amp;dispmax=20&amp;WebEnv=C%5B%60%3CIgCAA%5C%5Ee%3D%3FHCK%3CCDB%3CdlEjhCdbaiH%3F%40AARqjH%5EGe%5E%3EkDFH&amp;WebEnvRq=1</a> >; 3 pgs.
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<b>EXAMINER</b> 	<b>Date Considered</b> 12/30/03
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